## Amendments to the Claims

This listing of claims will replace all prior versions and listings of claims in the application:

- 1. (currently amended) A method for treating allergic rhinitis in mammals which comprises intranasally administering a pharmaceutically effective amount of a composition comprising an anti-allergy agent selected from the group consisting of emedastine and 0.01 0.8 % (w/v) olopatadine and 0.01 1.0 % (w/v) of a steroid selected from the group consisting of fluticasone, mometasone, budesonide and beclomethasone, wherein the composition has a pH of 3.5 8.0 and a viscosity of 1 50 cps.
- 2. (cancelled).
- 3. (cancelled).
- 4. (currently amended) The method of Claim 31 wherein the steroid is fluticasone.
- 5. (original) The method of Claim 1 wherein the steroid has an average particle size of  $2.5-5~\mu m$ .
- 6. (original) The method of Claim 1 wherein the steroid has an average particle size of less than 0.8 μm.
- 7. (original) The method of Claim 6 wherein the steroid has an average particle size of 0.5 µm or less.
- 8. (original) The method of Claim 1 wherein the composition is an aqueous composition packaged as a nasal spray.
- 9. (cancelled).

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10. (currently amended) A method for treating allergic rhinitis in mammals which comprises intranasally administering a pharmaceutically effective amount of a composition comprising 0.1 – 0.8 % (w/v) of olopatadine and 0.02 – 0.5 % (w/v) of a

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steroid selected from the group consisting of fluticasone, mometasone, budesonide and beclomethasone, wherein the composition has a pH of 3.5-8.0 and a viscosity of 1-50 cps., and the composition is an aqueous composition packaged as a nasal spray.